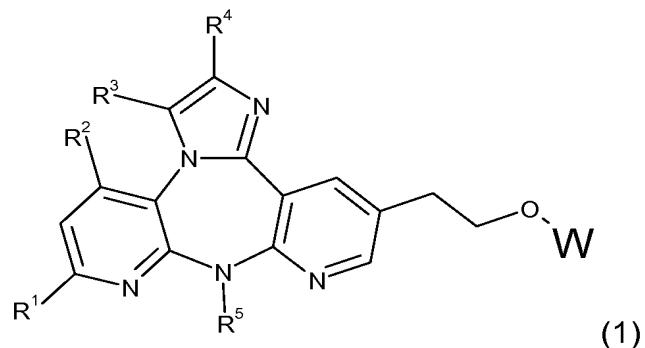


## **CLAIMS**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

Claim 1 (previously presented): A compound represented by formula 1:



wherein

**R<sup>1</sup>** is selected from the group consisting of H, halogen, (C<sub>1-4</sub>)alkyl, O(C<sub>1-4</sub>)alkyl, and haloalkyl;

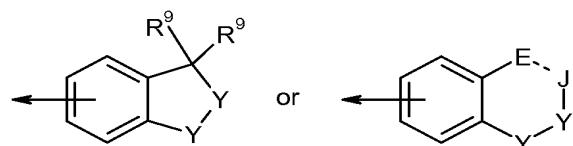
**R<sup>2</sup>** is H or Me;

**R<sup>3</sup>** is H or (C<sub>1-4</sub>)alkyl;

**R<sup>4</sup>** is H or (C<sub>1-4</sub>)alkyl;

**R<sup>5</sup>** is (C<sub>1-4</sub>)alkyl, (C<sub>1-4</sub>)alkyl(C<sub>3-7</sub>)cycloalkyl, or (C<sub>3-7</sub>)cycloalkyl; and

W is selected from:



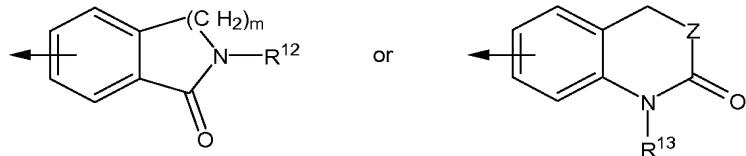
wherein.

a) one of **Y** is  $\text{SO}_2$  and the other **Y** is  $\text{NR}^6$ , provided that both are not the same, wherein  $\text{R}^6$  is selected from the group consisting of: H,  $\text{C}(\text{O})\text{O}(\text{C}_{1-4})\text{alkyl}$ ,  $(\text{C}_{1-4})\text{alkyl}$  or  $(\text{C}_{1-4})\text{alkyl}$  substituted with either a pyridinyl-N-oxide or  $\text{C}(\text{O})\text{OR}^8$  wherein  $\text{R}^8$  is H or  $(\text{C}_{1-4})\text{alkyl}$ ; and each  $\text{R}^9$  is independently H or  $(\text{C}_{1-4})\text{alkyl}$ ; and

b) **E** is  $CR^{10}R^{10}$  wherein each  $R^{10}$  is independently H or (C<sub>1-4</sub>) alkyl, **J** is CH<sub>2</sub> and the dotted line represents a single bond; or

c) **E** and **J** are both CR<sup>11</sup> wherein R<sup>11</sup> is H or (C<sub>1-4</sub>) alkyl and the dotted line represents a double bond; or

**W** is selected from:



wherein,

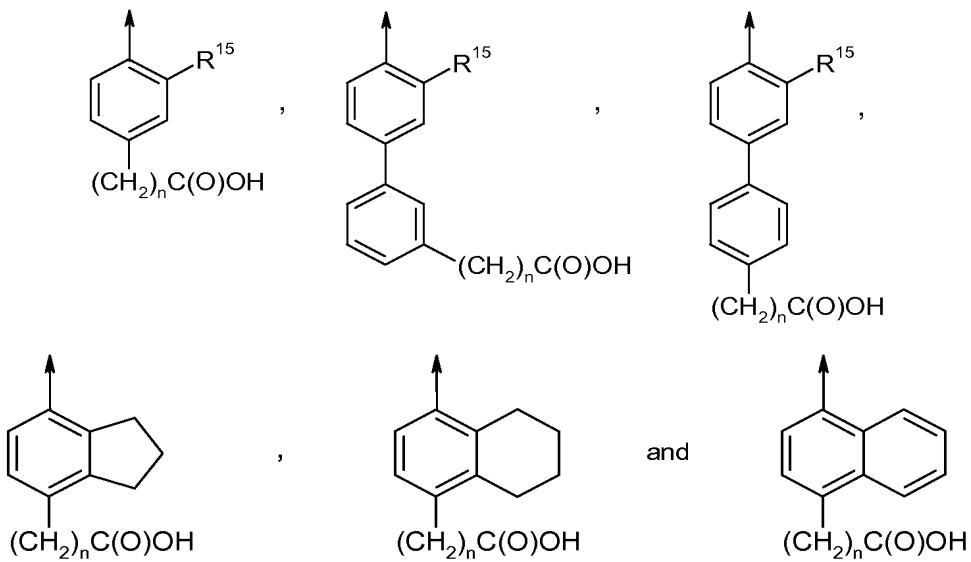
**m** is 1 or 2,

**R**<sup>12</sup> is H or C<sub>1-4</sub> alkyl,

**R**<sup>13</sup> is H or (C<sub>1-4</sub>) alkyl, and

**Z** is O or Z is NR<sup>14</sup> wherein R<sup>14</sup> is H or (C<sub>1-4</sub>) alkyl; or

**W** is selected from a group of aromatic radicals consisting of:

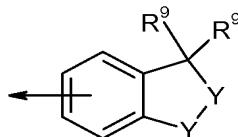


wherein **R**<sup>15</sup> is (C<sub>1-4</sub>) alkyl or CF<sub>3</sub>, and **n** is the integer 0, 1 or 2, or a pharmaceutically acceptable salt or ester thereof.

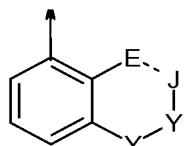
Claim 2 (original): The compound according to claim 1, wherein **R**<sup>1</sup> is selected from

the group consisting of: H, Cl, F, (C<sub>1-4</sub>) alkyl and CF<sub>3</sub>; R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> is each independently H or Me; R<sup>5</sup> is ethyl or cyclopropyl;

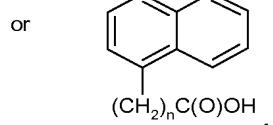
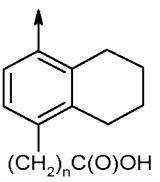
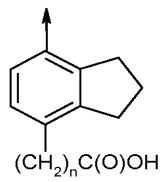
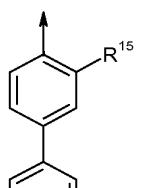
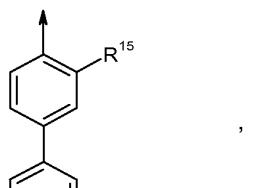
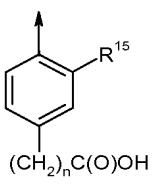
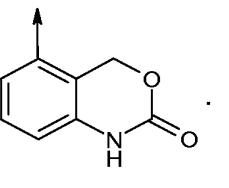
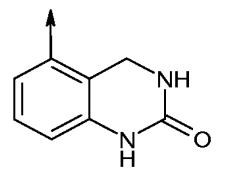
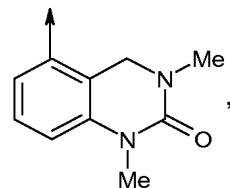
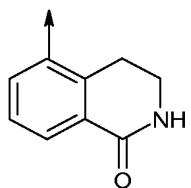
W is:



wherein Y is SO<sub>2</sub> and the other Y is NR<sup>6</sup>, provided that both are not the same, R<sup>6</sup> is H, C(O)OMe, C(O)OEt, (4-pyridinyl-N-oxide)methyl, CH<sub>2</sub>C(O)OH, CH<sub>2</sub>C(O)OMe, CH<sub>2</sub>C(O)OEt or CH<sub>2</sub>C(O)OCMe<sub>3</sub>, and each R<sup>9</sup> is independently H or Me; or



wherein E is CR<sup>10</sup>R<sup>10</sup> wherein each of R<sup>10</sup> is independently H or Me, J is CH<sub>2</sub> and the dotted line represents a single bond; or both E and J are CR<sup>11</sup> wherein R<sup>11</sup> is H or Me and the dotted line represents a double bond; one of Y is SO<sub>2</sub> and the other Y is NR<sup>6</sup> wherein R<sup>6</sup> is hydrogen or methyl; or

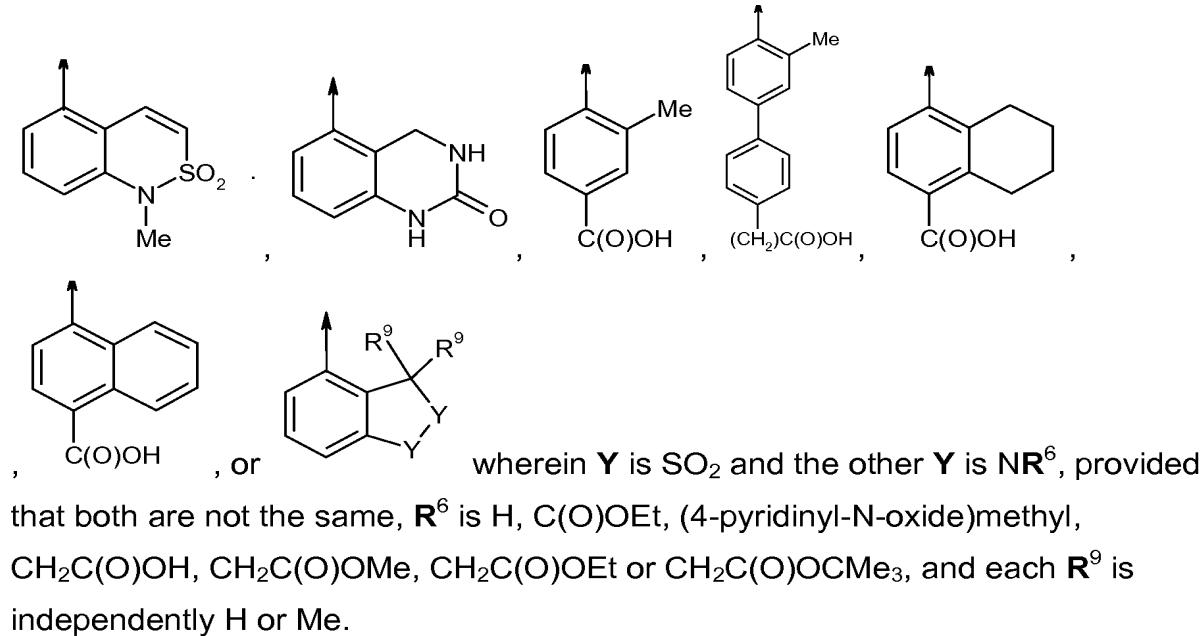


wherein R<sup>15</sup> is Me or Et, and n is 0 or 1.

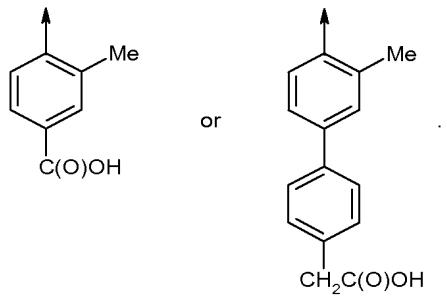
Claim 3 (original): The compound according to claim 2, wherein  $\mathbf{R}^{15}$  is Me.

Claim 4 (original): The compound according to claim 3, wherein  $\mathbf{R}^1$  is H, Cl, F and Me;  $\mathbf{R}^2$  is H or Me;

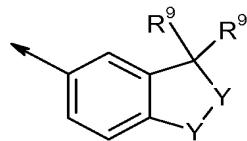
$\mathbf{W}$  is:



Claim 5 (original): The compound according to claim 4, wherein  $\mathbf{R}^3$  is Me,  $\mathbf{R}^6$  is H, C(O)OEt or (4-pyridinyl-N-oxide)methyl, and  $\mathbf{W}$  is:



Claim 6 (currently amended): The compound according to ~~claim 4~~ claim 3, wherein  $\mathbf{W}$  is:



wherein one **Y** is  $\text{SO}_2$  and the other **Y** is  $\text{NR}^6$ , provided that both are not the same,  $\text{R}^6$  is H,  $\text{C}(\text{O})\text{OEt}$ ,  $\text{CH}_2\text{C}(\text{O})\text{OH}$ ,  $\text{CH}_2\text{C}(\text{O})\text{OCMe}_3$ , (4-pyridinyl-N-oxide)methyl; and each  $\text{R}^9$  is independently H or Me.

Claim 7 (original): The compound according to claim 6, wherein  $\text{R}^6$  is H and each  $\text{R}^9$  is Me.

Claim 8 (cancelled)

Claim 9 (cancelled)

Claim 10 (cancelled)

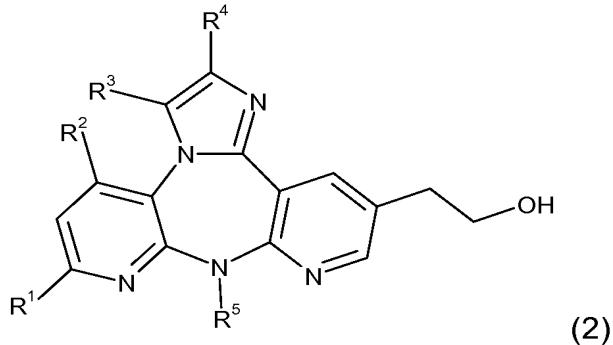
Claim 11 (previously presented): A pharmaceutical composition for the treatment of HIV infection, comprising a compound of formula 1 according to claim 1, or a pharmaceutically acceptable salt or ester thereof, in combination with a pharmaceutically acceptable carrier.

Claim 12 (previously presented): A method for the treatment of HIV infection, comprising administering to a patient an HIV inhibiting amount of a compound of formula 1 according to claim 1, or a pharmaceutically acceptable salt or ester thereof.

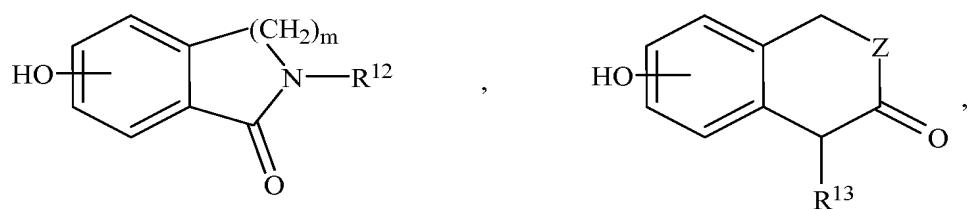
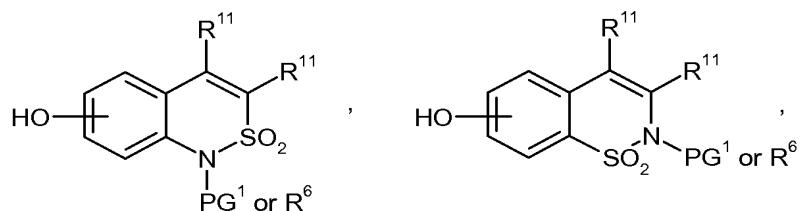
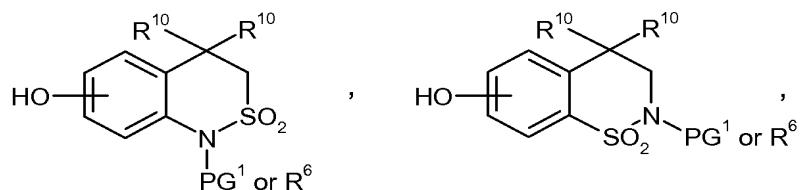
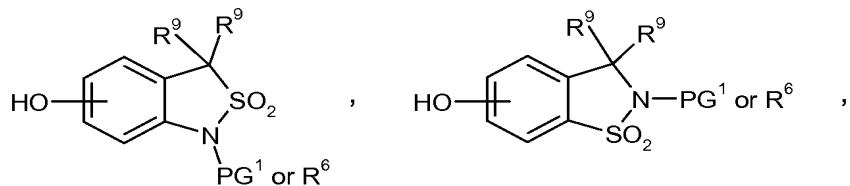
Claim 13 (previously presented): A method for the treatment of HIV infection, comprising administering to a patient an HIV inhibiting amount of a pharmaceutical composition according to claim 11.

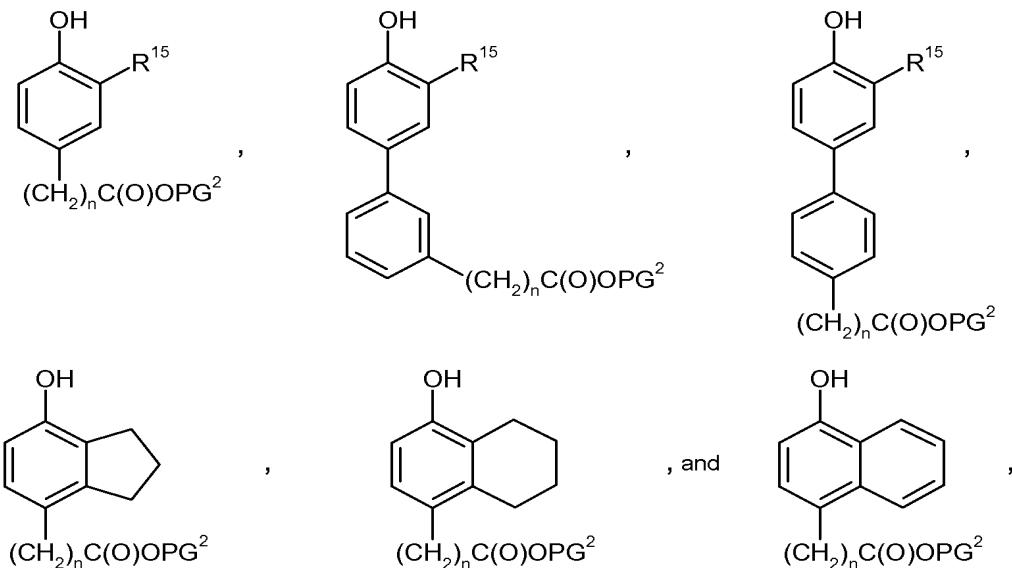
Claim 14 (previously presented): A process for producing a compound of formula 1 according to claim 1, comprising the step:

- coupling a compound of formula 2:



wherein **R<sup>1</sup>**, **R<sup>2</sup>**, **R<sup>3</sup>**, **R<sup>4</sup>**, and **R<sup>5</sup>** are as defined in claim 1, with a phenolic derivative selected from:



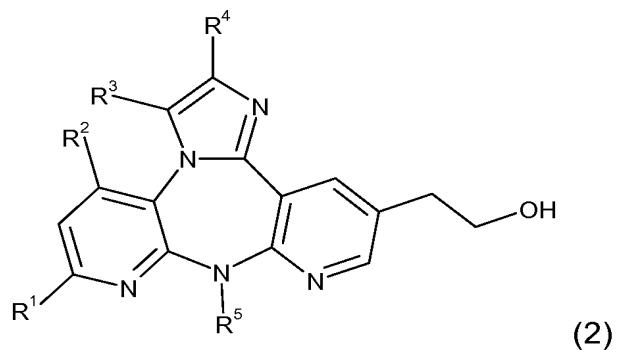


wherein PG<sup>1</sup> is a nitrogen protecting group and PG<sup>2</sup> is a carboxy protecting group, said protecting groups being removable under mildly acidic, mildly alkaline or reductive conditions, and R<sup>6</sup>, R<sup>9</sup>, R<sup>10</sup>, R<sup>11</sup>, R<sup>12</sup>, R<sup>13</sup>, R<sup>14</sup>, R<sup>15</sup>, m, n, and Z are as defined in claim 1.

Claim 15 (currently amended): The process according to claim 14, wherein said nitrogen protecting group carboxy protecting group is selected from: alkyl esters; aralkyl esters; and esters that can be cleaved by mild base treatment or mild reductive means.

Claim 16 (currently amended): The process according to claim 14, wherein said carboxy protecting group nitrogen protecting group is selected from: Boc (*tert*-butyloxycarbonyl) and alkyl carbamates.

Claim 17 (original): An intermediate compound of formula 2:



wherein **R**<sup>1</sup>, **R**<sup>2</sup>, **R**<sup>3</sup>, **R**<sup>4</sup>, and **R**<sup>5</sup> are as defined in claim 1.